

**IN THE UNITED STATES DISTRICT COURT
FOR THE DISTRICT OF DELAWARE**

PFIZER INC.,)	
PFIZER IRELAND PHARMACEUTICALS,)	
WARNER-LAMBERT COMPANY, and)	
WARNER-LAMBERT COMPANY LLC,)	
)	
Plaintiffs,)	
)	
v.)	C.A. No. 09-943-LPS and
)	C.A. No. 10-1135-LPS
DR. REDDY'S LABORATORIES LTD., and)	
DR. REDDY'S LABORATORIES INC.,)	
)	
Defendants.)	

**JOINT STIPULATION REGARDING CONSOLIDATION
AND AMENDMENT TO SCHEDULING ORDER**

Plaintiffs Pfizer Inc., Pfizer Ireland Pharmaceuticals, Warner-Lambert Company, and Warner-Lambert Company LLC (collectively "Pfizer") and Defendants Dr. Reddy's Laboratories Ltd. and Dr. Reddy's Laboratories Inc. (collectively "DRL"), through their respective attorneys, and subject to the approval of the Court, hereby agree and stipulate that:

1. This case, C.A. No. 09-943-LPS, is consolidated for all purposes with *Pfizer Inc. et al. v. Dr. Reddy's Labs. Ltd. et al.*, C.A. No. 10-1135-LPS. *All documents to be filed in 09-943-LPS.*
2. The Scheduling Order and Protective Order entered in C.A. No. 09-943-LPS shall govern both cases.
3. The Scheduling Order is hereby modified as follows and shall govern the consolidated action:

Stage of Discovery	Proposed Completion Date
Substantial completion of document production ¹	March 16, 2011
Completion of supplemental production (if any)	April 1, 2011
Completion of fact discovery	May 1, 2011
Opening Expert Reports Exchanged	May 16, 2011
Rebuttal Expert Reports Exchanged	June 30, 2011
Reply Expert Reports Exchanged	July 14, 2011
Completion of expert discovery	July 28, 2011
Deadline for filing case dispositive motions	August 11, 2011
Deadline for filing responses to case dispositive motions	September 1, 2011
Deadline for filing replies to case dispositive motions	September 15, 2011
Trial	December 5, 2011

Dated: March 2, 2011

/s/ Mary W. Bourke

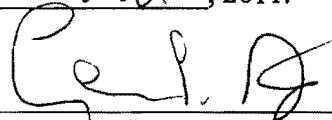
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Attorneys for Plaintiffs

SO ORDERED this 3rd day of March, 2011.


UNITED STATES DISTRICT JUDGE

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¹ This includes all basic documents for the 80 mg ANDA tablet, i.e., the ANDA and DMF, XRPD test results that can be produced by this date, batch records, and other regulatory documents.